In the claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1. (Canceled).
- (Currently Amended) A method of providing hematopoietic stem cells to a subject comprising the steps of:

administering a thrombopoietin (TPO) mimetic compound to a subject to increase stem cells in said subject;

harvesting one or more of the stem cells;

treating said subject with a bone marrow ablative treatment; and transplanting the harvested stem cells into the subject.

wherein the TPO mimetic compound has the following sequence:

wherein (2-Nal) is β-(2-naphthyl)alanine and X10-is Sar is sarcosine.

- 3. (Previously Presented) The method of claim 2, wherein the subject is a human.
- (Original) The method of claim 2, wherein the one or more stem cells are cryopreserved after harvesting.
- 5. (Original) The method of claim 4, wherein the one or more cryopreserved stem cells are thawed and determined to be viable prior to transplanting the stem cells into the subject.
- (Original) The method of claim 4, wherein the one or more stem cells are transplanted into the subject when the subject is in need of such transplantation.
- 7. (Previously Presented) The method of claim 2, wherein the TPO mimetic compound

has reduced immunogenicity relative to one or more of rhTPO and rhIL-11.

- 8. (Previously Presented) The method of claim 2, wherein the TPO mimetic compound has an improved pharmacokinetic profile relative to one or more of rhTPO and rhIL-11.
- (Currently Amended) A method of reducing a time to engraftment following reinfusion of stem cells in a subject comprising the steps of:

administering a thrombopoietin (TPO) mimetic compound to the subject;

increasing stem cells in said subject;

harvesting one or more of the stem cells;

treating said subject with a bone marrow ablative treatment; and

transplanting the one or more harvested stem cells into the subject,

wherein the TPO mimetic compound has the following sequence:

IEGPTLRO(2-Nal)LAAR-(Sar)

K(NH2)

IEGPTLRQ(2-Nal)LAAR-(Sar),

wherein (2-Nal) is β-(2-naphthyl)alanine and X10-is Sar is sarcosine.

10. (Currently Amended) A method of reducing the incidence of delayed primary engraftment comprising the steps of:

administering a thrombopoietin (TPO) mimetic compound to the subject:

increasing stem cells in said subject:

harvesting one or more of the stem cells:

treating said subject with a bone marrow ablative treatment; and

transplanting the one or more harvested stem cells into the subject,

wherein the TPO mimetic compound has the following sequence:

IEGPTLRQ(2-Nal)LAAR-(Sar)

K(NH2)

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LEGPTLRO(2-Nal)LAAR-(Sar).

wherein (2-NaI) is β-(2-naphthyl)alanine and X-10-is Sar is sarcosine.

11. (Currently Amended) A method of reducing the incidence of secondary failure of platelet production comprising the steps of:

administering a thrombopoietin (TPO) mimetic compound to the subject;

increasing stem cells in said subject;

harvesting one or more the stem cells;

treating the subject with a bone marrow ablative treatment; and

transplanting the one or more harvested stem cells into the subject,

wherein the TPO mimetic compound has the following sequence:

\ K(NH₂)

IEGPTLRO(2-Nal)LAAR-(Sar),

wherein (2-Nal) is β-(2-naphthyl)alanine and X10-is Sar is sarcosine.

12. (Currently Amended) A method of reducing the time of platelet and/or neutrophil engraftment following reinfusion of stem cells in a subject comprising the steps of:

administering a thrombopoietin (TPO) mimetic compound to the subject;

increasing stem cells in said subject:

harvesting one or more of the stem cells;

treating the subject with a bone marrow ablative treatment; and

transplanting the one or more harvested stem cells into the subject,

wherein the TPO mimetic compound has the following sequence:

K(NH2)

IEGPTLRO(2-Nal)LAAR-(Sar).

wherein (2-Nal) is β-(2-naphthyl)alanine and X+++is Sar is sarcosine.

| 13. Canceled. |
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| 14. Canceled. |
| 15. (Previously Presented) The method of claim 2, wherein said TPO mimetic compound i covalently attached to a hydrophilic polymer. |
| 16. (Previously Presented) The method of claim 15, wherein said hydrophilic polymer has an average molecular weight of between about 500 to about 40,000 daltons. |
| 17. (Previously Presented) The method of claim 16, wherein said hydrophilic polymer has an average molecular weight of between about 5,000 to about 20,000 daltons. |
| $18. \ (Previously\ Presented)\ The\ method\ of\ claim\ 17,\ wherein\ said\ hydrophilic\ polymer\ has\ an\ average\ molecular\ weight\ of\ about\ 20,000\ daltons.$ |
| 19. (Previously Presented) The method of claim 15, wherein said polymer is polyethylene glycol. |
| 20. Canceled. |
| 21. Canceled. |
| 22. Canceled. |
| 23. Canceled. |
| 24. Canceled. |
| 25. Canceled. |
| 26. (Previously Presented) The method of claim 2, wherein each of the dimeric subunits |

of said TPO mimetic compound is covalently attached to a hydrophilic polymer.

- 27. (Previously Presented) The method of claim 26, wherein said hydrophilic polymer has an average molecular weight of between about 500 to about 40,000 daltons.
- 28. (Previously Presented) The method of claim 27, wherein said hydrophilic polymer has an average molecular weight of between about 5,000 to about 20,000 daltons.
- 29. (Previously Presented) The method of claim 28, wherein said hydrophilic polymer has an average molecular weight of about 20,000 daltons.
- (Previously Presented) The method of claim 26, wherein said polymer is polyethylene glycol.
- 31. (Previously Presented) The method of claim 2, wherein said stem cells are within said subject's bone marrow.
- 32. (Previously Presented) The method of claim 2, wherein said stem cells are within said subject's peripheral circulation.
- 33. (Previously Presented) The method of claim 2, wherein said subject is treated with chemotherapy.
- 34. (Previously Presented) The method of claim 2, wherein said subject is treated with radiation thereapy.